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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/082,476	02/20/2002	Gregory D. May	NAPRO-3	4408
25213	7590	01/18/2005	EXAMINER	
HELLER EHRMAN WHITE & MCAULIFFE LLP			FREDMAN, JEFFREY NORMAN	
275 MIDDLEFIELD ROAD			ART UNIT	
MENLO PARK, CA 94025-3506			PAPER NUMBER	
			1637	

DATE MAILED: 01/18/2005

Please find below and/or attached an Office communication concerning this application or proceeding.

**Office Action Summary**

Application No.

10/082,476

Applicant(s)

MAY ET AL.

Examiner

Jeffrey Fredman

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

**Period for Reply**

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

**Status**

- 1) ☒ Responsive to communication(s) filed on 29 November 2004.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

**Disposition of Claims**

- 4) ☒ Claim(s) 13-17 and 29-32 is/are pending in the application.
- 4a) Of the above claim(s) \_\_\_\_\_ is/are withdrawn from consideration.
- 5) ☐ Claim(s) \_\_\_\_\_ is/are allowed.
- 6) ☒ Claim(s) 13-17 and 29-32 is/are rejected.
- 7) ☐ Claim(s) \_\_\_\_\_ is/are objected to.
- 8) ☐ Claim(s) \_\_\_\_\_ are subject to restriction and/or election requirement.

**Application Papers**

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on \_\_\_\_\_ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.  
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).  
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

**Priority under 35 U.S.C. § 119**

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some \* c) ☐ None of:
- ☐ Certified copies of the priority documents have been received.
  - ☐ Certified copies of the priority documents have been received in Application No. \_\_\_\_\_.
  - ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

\* See the attached detailed Office action for a list of the certified copies not received.

**Attachment(s)**

- 1) ☐ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☒ Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08)  
Paper No(s)/Mail Date 7/06/04; 2/17/04
- 4) ☐ Interview Summary (PTO-413)  
Paper No(s)/Mail Date. \_\_\_\_\_
- 5) ☐ Notice of Informal Patent Application (PTO-152)
- 6) ☐ Other: \_\_\_\_\_

## **DETAILED ACTION**

### ***Continued Examination Under 37 CFR 1.114***

1. A request for continued examination under 37 CFR 1.114, including the fee set forth in 37 CFR 1.17(e), was filed in this application after final rejection. Since this application is eligible for continued examination under 37 CFR 1.114, and the fee set forth in 37 CFR 1.17(e) has been timely paid, the finality of the previous Office action has been withdrawn pursuant to 37 CFR 1.114. Applicant's submission filed on November 29, 2004 has been entered.

### ***Claim Rejections - 35 USC § 103***

2. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

This application currently names joint inventors. In considering patentability of the claims under 35 U.S.C. 103(a), the examiner presumes that the subject matter of the various claims was commonly owned at the time any inventions covered therein were made absent any evidence to the contrary. Applicant is advised of the obligation under 37 CFR 1.56 to point out the inventor and invention dates of each claim that was not commonly owned at the time a later invention was made in order for the examiner to consider the applicability of 35 U.S.C. 103(c) and potential 35 U.S.C. 102(e), (f) or (g) prior art under 35 U.S.C. 103(a).

3. Claims 13-17 and 29-32 are rejected under 35 U.S.C. 103(a) as being unpatentable over Yamashita et al (EP 718,404 A2, June 1996) in view of Baszcynski et al (U.S. 6,528,700).

Yamashita teaches a composition comprising:

- a) a duplex DNA comprising the lacZ (see page 4, example 1 where LacZ is used and page 5, table I, the PUC-Mut solution).
- b) an oligonucleotide capable of introducing a site specific predetermined change in the LacZ target sequence (see page 5, table I, here the dR1 solution)
- c) a cell free extract (see page 5, table I, here the RecA protein solution)
- d) a reaction buffer (see page 5, table I, here the 10x reaction buffer).

With regard to claims 14-15, Yamashita teaches an oligonucleotides that are approximately 22-24 nucleotides in length (see SEQ ID Nos: 1-3).

With regard to claim 16, Yamashita teaches an oligonucleotide with a single 3' and 5' end (see SEQ ID NO: 1).

With regard to claims 17 and 30, Yamashita teaches the LacZ gene, which is linked to a promoter that can be expressed (see page 4, lines 15-25).

With regard to claims 31 and 32, Yamashita teaches the duplex DNA is a plasmid DNA (see page 4, lines 18-25).

Yamashita does not teach the use of a plant cell extract or the oligonucleotide of claim 29.

Baszcynski teaches a composition for gene correction comprising:

- a) a duplex DNA (see column 14, lines 60-65).
- b) an oligonucleotide (see column 14, lines 60-65).
- c) a plant cell free extract (see column 14, lines 60-65).
- d) a reaction buffer (see column 14, lines 60-65).

With regard to claims 14-15, Baszcynski teaches an oligonucleotide that is approximately 90 nucleotides in length (see SEQ ID NO: 2).

With regard to claim 16, Baszcynski teaches an oligonucleotide with a single 3' and 5' end (see SEQ ID NO: 2).

With regard to claims 17 and 30, Baszcynski teaches the double stranded DNA including the pPHPP10247 plasmid which comprises the AHAS gene under the control of the ubiquitin promoter (see column 11, lines 49-67).

With regard to claim 29, Baszcynski teaches a self complementary oligonucleotide with at least 5 bases that are base paired (see figure 7).

It would have been prima facie obvious to one of ordinary skill in the art at the time the invention was made to use a plant cell extract as taught by Baszcynski in the composition of Yamashita since Yamashita notes "Namely, proteins being similar to RecA but originating in other sources and variants thereof are usable herein (see page 3, lines 5-7)." Yamashita further recognizes the equivalence of plants and other cell types, noting that host cells can include "bacteria such as Escherichia coli and Bacillus subtilis, yeasts, fungi, plant cells and animal cells (see page 3, line 21)." Therefore an

ordinary practitioner would have been motivated by Yamashita to use other cell free extracts from sources which would function in gene correction, including equivalent extracts from plants. Baszcynski motivates the use of plant extracts since Baszcynski teaches that "Compositions and methods for targeted gene correction, conversion, or modification in plants are provided (see column 4, lines 66-67)." So an ordinary practitioner would have been motivated by Baszcynski to use plants for the modification since Baszcynski teaches in vitro plant cell extracts (see column 60) and wants to gene correct plants (see abstract), while Yamashita recognizes the equivalence and use of other cell free extracts.

A further motivation is that of substituting equivalents, where the plant cell free extract is an equivalent, since Yamashita recognizes the equivalence of the different host cell types, as MPEP 2144.06 notes " Substituting equivalents known for the same purpose. In order to rely on equivalence as a rationale supporting an obviousness rejection, the equivalency must be recognized in the prior art, and cannot be based on applicant's disclosure or the mere fact that the components at issue are functional or mechanical equivalents. An express suggestion to substitute one equivalent component or process for another is not necessary to render such substitution obvious. In re Fout , 675 F.2d 297, 213 USPQ 532 (CCPA 1982)."

Further, an ordinary practitioner would have been further motivated to use oligonucleotides with self base pairing since Baszcynski notes that these oligonucleotides are resistant to nuclease digestion and present no impediment to

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pairing with the target, thereby improving the sensitivity and specificity of the assay (see column 5, lines 31-38).

The ordinary practitioner would have had a reasonable expectation of success given that Baszcynski demonstrated that in vivo gene correction in plants functioned (see column 20) and since Yamashita demonstrated successful in vitro gene correction (see example 3).

### ***Response to Arguments***

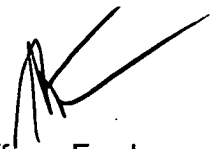
4. Applicant's arguments with respect to the claims have been considered but are moot in view of the new ground(s) of rejection.

### ***Conclusion***

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Jeffrey Fredman whose telephone number is (571)272-0742. The examiner can normally be reached on 6:30-4:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Gary Benzion can be reached on (571)272-0782. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).



Jeffrey Fredman  
Primary Examiner  
Art Unit 1637

